Applicant: Vanda A. Lennon et al. Attorney's Docket No.: 07039-497001 / MMV-03-132

Serial No.: 10/723,180

Filed: November 25, 2003

Page : 2 of 4

Amendments to the Claims:

Please amend claims 1 and 2 as follows. Please cancel claims 3 and 7-17 without prejudice to continued prosecution. The claims and their status are shown below.

1. (Currently Amended) A method of detecting the presence or absence of a neuromyelitis optica (NMO) [[NMO]]-specific autoantibody in a biological sample from an individual, comprising the steps of:

contacting said biological sample with a NMO antigenic polypeptide or fragment thereof, wherein said NMO antigenic polypeptide is aquaporin-4, wherein said fragment of said NMO antigenic polypeptide is a polypeptide for which a NMO-specific autoantibody has specific binding affinity; and

detecting the presence or absence of binding of said NMO antigenic polypeptide to said NMO-specific autoantibody in said biological sample,

wherein the presence of said binding of said NMO antigenic polypeptide to said NMO-specific autoantibody is indicative of NMO in said individual.

- 2. (Currently Amended) The method of claim 1, wherein the presence of said NMO-specific autoantibody in said biological sample is associated with vision impairment, weakness, numbness, spasms or abnormal or painful sensations, and/or loss of bladder control, and/or or loss of bowel control in said individual.
 - 3. (Canceled)
- 4. (Original) The method of claim 1, wherein said NMO antigenic polypeptide is a recombinantly-expressed NMO antigenic polypeptide.
- 5. (Original) The method of claim 1, wherein said NMO-specific polypeptide is in a solid tissue selected from the group consisting of brain, spinal cord, optic nerve, kidney, or stomach.
- 6. (Original) The method of claim 1, wherein said biological sample is selected from the group consisting of blood, serum, plasma, and cerebrospinal fluid.

7-17. (Canceled)